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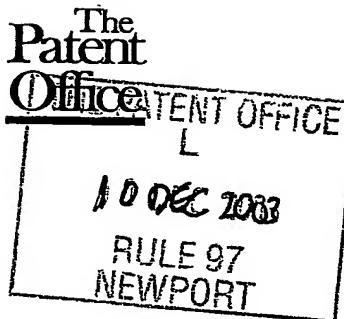
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1. Your reference

P103734GB

2. Patent application number

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3. Full name, address and postcode of the or of each applicant (underline all surnames)

Ludwig Institute for Cancer Research
Postfach 8024
Zurich
Switzerland

Patents ADP number (if you know it)

577155002

If the applicant is a corporate body, give the country/state of its incorporation

CH

4. Title of the invention

Tumour Suppressor Protein

5. Name of your agent (if you have one)

Harrison Goddard Foote

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

31 St Saviourgate
YORK
YO1 8NQ

Patents ADP number (if you know it)

07914237002

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Country

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Rob Docherty

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Tumour Suppressor Protein

The invention relates to a protein which binds and modulates the activity of a tumour suppressor protein, for example p53; a nucleic acid molecule encoding 5 said protein and screening methods which modulate the binding activity of said polypeptide for its target polypeptide.

Tumour suppressor genes encode proteins which function to inhibit cell growth or division and are therefore important with respect to maintaining 10 proliferation, growth and differentiation of normal cells. Mutations in tumour suppressor genes result in abnormal cell-cycle progression whereby the normal cell-cycle check points which arrest the cell-cycle, when, for example, DNA is damaged, are ignored and damaged cells divide uncontrollably. The products of tumour suppressor genes function in all parts of the cell (e.g. cell surface, 15 cytoplasm, nucleus) to prevent the passage of damaged cells through the cell-cycle (i.e. G1, S, G2, M and cytokinesis). A number of tumour suppressor genes have been isolated and sequenced. These include the Retinoblastoma gene (Rb), mutations in which are linked to cancers such as bone (osteocarcoma), bladder, small cell lung and breast cancer, as well as 20 retinoblastoma. The Wilms Tumour 1 gene (WT-1), mutations in which are linked to nephroblastoma and neurofibromatosis.

Arguably the tumour suppressor gene which has been the subject of the most intense research is p53. p53 encodes a protein which functions as a 25 transcription factor and is a key regulator of the cell division cycle. It was discovered in 1978 (Lane and Crawford, 1979) as a protein shown to bind with affinity to the SV40 large T antigen. The p53 gene encodes a 393 amino acid polypeptide with a molecular weight of 53kDa. One of the most important tumour suppression functions of p53 is its ability to induce apoptosis

Apoptosis, or programmed cell death, is a process by which multi-cellular organisms regulate cell number and differentiation. The process is regulated by factors which either induce or prevent apoptosis. Inducers of apoptosis include Bcl-2 family members, caspase family members and their associated factors

5 Apaf-1 and Fadd. Caspases are synthesised as proenzymes which become activated after proteolytic cleavage. The active caspase then induces many of the morphological and biochemical changes associated with apoptosis. Mitochondria play a pivotal role in the activation process through the release of pro-apoptotic factors such as cytochrome c, AIF and Diablo. The release from

10 mitochondria is controlled by the Bcl-2 family of proteins; (e.g. Bcl-2 and Bcl-xL inhibit release; Bax and Bak induce release).

The polypeptide referred to as iASPP that is described in WO02/12325 is a further example of an agent involved in the regulation of apoptosis.

15

We describe a variant iASPP polypeptide which has characteristics which are distinct from those described in W02/12325. The polypeptide, referred to as iASPP 6C, is extended at its amino terminus by **X amino acid residues** and binds preferentially to p53 when compared to iASPP. iASPP C6 preferentially binds p53 when compared to the shorter version described in W02/12325. The shorter version preferentially binds the apoptosis inducer protein Bcl 2.

iASPP C6 is a ubiquitinated polypeptide which likely controls the turnover of iASPP C6 *in vivo*. Ubiquitin is a small protein made up of 76 amino acids

25 which is highly conserved across species. The most important function assigned to ubiquitin is in regulating protein turnover. Research in recent years has identified many accessory proteins involved in ubiquitin induced proteolysis. The first step is the ligation of ubiquitin to a target protein which is destined for degradation. This is mediated by three proteins referred to as E1,

30 E2 and E3. Ubiquitin is first activated by E1 activating enzyme, a homodimer composed of two identical 105kDa subunits which is ligated to ubiquitin via a

thioester bond. Following activation the E1:ubiquitin conjugate is transported by E2 (referred to as a carrier protein). The E2 proteins vary markedly in size but do have some conserved elements. The E2 protein accepts the ubiquitin from E1 and forms a second complex again via a thioester bond. The E3 5 protein may or may not become involved in the final step, which is the transfer of ubiquitin to a protein substrate. This is followed by recognition by a protease, which degrades the ubiquitinated protein. The protease may be part of a structure referred to as the proteosome which is a large multi-subunit complex of proteases and associated co-factors. In some examples proteins can 10 become polyubiquitinated, which results from ubiquitin proteins being ligated to ubiquitin proteins, which are already ligated to a target protein.

According to an aspect of the invention there is provided an isolated polypeptide wherein said polypeptide is represented by the amino acid 15 sequence as shown in Figure 1a, or a variant polypeptide which variant is modified by addition, deletion or substitution of at least one amino acid residue characterised in that said polypeptide has the following characteristics:

- i) a polypeptide which preferentially binds the tumour suppressor polypeptide p53 to inhibit the pro-apoptotic activity of p53 when compared to a polypeptide, or variant thereof, as represented by the amino acid sequence as shown in Figure 2a;
- ii) a polypeptide which includes at least one amino acid residue which residue is ubiquitinated; and
- iii) a polypeptide which comprises an amino-terminal polypeptide domain wherein said domain is represented between amino acid 1 and 483 of the amino acid sequence shown in Figure 1a.

In a preferred embodiment of the invention said polypeptide preferentially binds p53 when compared to a polypeptide represented by the amino acid 30 sequence shown in Figure 2a.

In a further preferred embodiment of the invention said polypeptide is modified by addition, deletion or substitution of at least one amino acid residue wherein said modification is between amino acid residues +1 and +483 of the amino acid sequence presented in Figure 1a.

5

Assays to determine the binding of polypeptides, which are herein disclosed, to for example, p53 are known in the art and described in the present application.

In a further preferred embodiment of the invention said polypeptide comprises 10 the amino acid sequence shown in Figure 1a. Preferably said polypeptide consists of the amino acid sequence shown in Figure 1a.

15 According to an aspect of the invention there is provided an isolated nucleic acid molecule wherein said nucleic acid molecule encodes a polypeptide according to the invention.

In a preferred embodiment of the invention said nucleic acid molecule is represented by the nucleic acid sequence shown in Figure 1b or a nucleic acid molecule which hybridises to the sequence shown in Figure 1b under stringent 20 hybridisation conditions and which encodes a polypeptide according to the invention.

In a preferred embodiment of the invention said nucleic acid molecule consists 25 of the nucleic acid sequence shown in Figure 1b.

In a further preferred embodiment of the invention said isolated nucleic acid molecule is a cDNA. In an alternative preferred embodiment of the invention said nucleic acid molecule is genomic DNA.

30 According to a further aspect of the invention there is provided a vector comprising a nucleic acid molecule according to the invention. Preferably said

vector is an expression vector adapted for recombinant expression of said polypeptide.

5 Preferably, said vector is adapted for prokaryotic gene expression. In an alternative embodiment of the invention said vector is adapted for eukaryotic gene expression.

10 Typically said adaptation includes, by example and not by way of limitation, the provision of transcription control sequences (promoter sequences) which mediate cell/tissue specific expression. These promoter sequences may be cell/tissue specific, inducible or constitutive.

15 Promoter is an art recognised term and includes the following features which are provided by example only, and not by way of limitation. Enhancer elements are *cis* acting nucleic acid sequences often found 5' to the transcription initiation site of a gene (enhancers can also be found 3' to a gene sequence or even located in intronic sequences and is therefore position independent). Enhancers function to increase the rate of transcription of the gene to which the enhancer is linked. Enhancer activity is responsive to *trans* acting transcription 20 factors (polypeptides) which have been shown to bind specifically to enhancer elements. The binding/activity of transcription factors (please see Eukaryotic Transcription Factors, by David S Latchman, Academic Press Ltd, San Diego) is responsive to a number of environmental cues which include, by example and not by way of limitation, intermediary metabolites or environmental 25 effectors, for example temperature.

Promoter elements also include so-called TATA box and RNA polymerase initiation selection (RIS) sequences which function to select a site of transcription initiation. These sequences also bind polypeptides which function, 30 *inter alia*, to facilitate transcription initiation selection by RNA polymerase.

Adaptations also include the provision of selectable markers and autonomous replication sequences which both facilitate the maintenance of said vector in either the eukaryotic cell or prokaryotic host. Vectors which are maintained autonomously are referred to as episomal vectors. Episomal vectors are desirable since these molecules can incorporate large DNA fragments (30-50kb DNA). Episomal vectors of this type are described in WO98/07876.

Adaptations which facilitate the expression of vector encoded genes include the provision of transcription termination/polyadenylation sequences. This also includes the provision of internal ribosome entry sites (IRES) which function to maximise expression of vector encoded genes arranged in bicistronic or multi-cistronic expression cassettes.

These adaptations are well known in the art. There is a significant amount of published literature with respect to expression vector construction and recombinant DNA techniques in general. Please see, Sambrook et al (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbour Laboratory, Cold Spring Harbour, NY and references therein; Marston, F (1987) DNA Cloning Techniques: A Practical Approach Vol III IRL Press, Oxford UK; DNA Cloning: F M Ausubel et al, Current Protocols in Molecular Biology, John Wiley & Sons, Inc.(1994).

According to a fourth aspect of the invention there is provided a method for the production of the polypeptide according to the invention comprising:

- 25 i) providing a cell transformed/transfected with a nucleic acid molecule according to the invention;
- ii) growing said cell in conditions conducive to the manufacture of said polypeptide; and
- i) purifying said polypeptide from said cell, or its growth environment.

In a preferred embodiment of the invention said nucleic acid molecule is the vector according to the invention.

In a preferred method of the invention said vector encodes, and thus said recombinant polypeptide is provided with, a secretion signal to facilitate 5 purification of said polypeptide.

According to a further aspect of the invention there is provided an antibody which binds the polypeptide according to the invention characterised in that said antibody binds said polypeptide between amino acid residues +1 to +483 10 of the amino acid sequence shown in Figure 1a.

Preferably said antibody does not bind said polypeptide represented by the sequence +484 to +828 of the amino acid sequence shown in Figure 1a.

15 Antibodies, also known as immunoglobulins, are protein molecules which usually have specificity for foreign molecules (antigens). Immunoglobulins (Ig) are a class of structurally related proteins consisting of two pairs of polypeptide chains, one pair of light (L) (low molecular weight) chain (κ or λ), and one pair of heavy (H) chains (γ , α , μ , δ and ϵ), all four linked together by 20 disulphide bonds. Both H and L chains have regions that contribute to the binding of antigen and that are highly variable from one Ig molecule to another. In addition, H and L chains contain regions that are non-variable or constant.

25 The L chains consist of two domains. The carboxy-terminal domain is essentially identical among L chains of a given type and is referred to as the “constant” (C) region. The amino terminal domain varies from L chain to L chain and contributes to the binding site of the antibody. Because of its variability, it is referred to as the “variable” (V) region.

30 The H chains of Ig molecules are of several classes, α , μ , σ , α , and γ (of which there are several sub-classes). An assembled Ig molecule consisting of one or

more units of two identical H and L chains, derives its name from the H chain that it possesses. Thus, there are five Ig isotypes: IgA, IgM, IgD, IgE and IgG (with four sub-classes based on the differences in the 'constant' regions of the H chains, i.e., IgG1, IgG2, IgG3 and IgG4). Further detail regarding antibody structure and their various functions can be found in, Using Antibodies: A laboratory manual, Cold Spring Harbour Laboratory Press.

In a preferred embodiment of the invention said fragment is a Fab fragment.

10 In a further preferred embodiment of the invention said antibody is selected from the group consisting of: F(ab')₂, Fab, Fv and Fd fragments; and antibodies comprising CDR3 regions.

15 Preferably said fragments are single chain antibody variable regions (scFV's) or domain antibodies. If a hybridoma exists for a specific monoclonal antibody it is well within the knowledge of the skilled person to isolate scFv's from mRNA extracted from said hybridoma via RT PCR. Alternatively, phage display screening can be undertaken to identify clones expressing scFv's. Domain antibodies are the smallest binding part of an antibody (approximately 13kDa).
20 Examples of this technology is disclosed in US6, 248, 516, US6, 291, 158, US6,127, 197 and EP0368684 which are all incorporated by reference in their entirety.

25 A modified antibody, or variant antibody and reference antibody, may differ in amino acid sequence by one or more substitutions, additions, deletions, truncations which may be present in any combination. Among preferred variants are those that vary from a reference polypeptide by conservative amino acid substitutions. Such substitutions are those that substitute a given amino acid by another amino acid of like characteristics. The following non-limiting 30 list of amino acids are considered conservative replacements (similar): a) alanine, serine, and threonine; b) glutamic acid and aspartic acid; c)

asparagine and glutamine d) arginine and lysine; e) isoleucine, leucine, methionine and valine and f) phenylalanine, tyrosine and tryptophan. Most highly preferred are variants which show enhanced biological activity.

5 Preferably said antibody is a humanised or chimeric antibody.

A chimeric antibody is produced by recombinant methods to contain the variable region of an antibody with an invariant or constant region of a human antibody.

10

A humanised antibody is produced by recombinant methods to combine the complementarity determining regions (CDRs) of an antibody with both the constant (C) regions and the framework regions from the variable (V) regions of a human antibody.

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Chimeric antibodies are recombinant antibodies in which all of the V-regions of a mouse or rat antibody are combined with human antibody C-regions. Humanised antibodies are recombinant hybrid antibodies which fuse the complementarity determining regions from a rodent antibody V-region with the

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framework regions from the human antibody V-regions. The C-regions from the human antibody are also used. The complementarity determining regions (CDRs) are the regions within the N-terminal domain of both the heavy and light chain of the antibody to where the majority of the variation of the V-region is restricted. These regions form loops at the surface of the antibody molecule. These loops provide the binding surface between the antibody and antigen.

25

Antibodies from non-human animals provoke an immune response to the foreign antibody and its removal from the circulation. Both chimeric and humanised antibodies have reduced antigenicity when injected to a human subject because there is a reduced amount of rodent (i.e. foreign) antibody

within the recombinant hybrid antibody, while the human antibody regions do not elicit an immune response. This results in a weaker immune response and a decrease in the clearance of the antibody. This is clearly desirable when using therapeutic antibodies in the treatment of human diseases. Humanised 5 antibodies are designed to have less "foreign" antibody regions and are therefore thought to be less immunogenic than chimeric antibodies.

According to a further aspect of the invention there is provided a polypeptide according to the invention for use as a pharmaceutical.

10

According to a further aspect of the invention there is provided a nucleic acid according to the invention for use as a pharmaceutical.

In a preferred embodiment of the invention said pharmaceutical further comprises a diluent, carrier or excipient.

15

When administered, the therapeutic compositions of the present invention are administered in pharmaceutically acceptable preparations. Such preparations may routinely contain pharmaceutically acceptable concentrations of salt, buffering agents, preservatives, compatible carriers, supplementary immune 20 potentiating agents such as adjuvants and cytokines and optionally other therapeutic agents, such as chemotherapeutic agents.

The therapeutics of the invention can be administered by any conventional route, including injection or by gradual infusion over time. The administration 25 may, for example, be oral, intravenous, intraperitoneal, intramuscular, intracavity, subcutaneous, or transdermal. When antibodies are used therapeutically, a preferred route of administration is by pulmonary aerosol. Techniques for preparing aerosol delivery systems containing antibodies are well known to those of skill in the art. Generally, such systems should utilize 30 components which will not significantly impair the biological properties of the

antibodies, such as the paratope binding capacity (see, for example, Sciarra and Cutie, "Aerosols," in Remington's Pharmaceutical Sciences, 18th edition, 1990, pp 1694-1712; incorporated by reference). Those of skill in the art can readily determine the various parameters and conditions for producing antibody aerosols without resort to undue experimentation. When using antisense preparations of the invention, slow intravenous administration is preferred.

The compositions of the invention are administered in effective amounts. An "effective amount" is that amount of a composition that alone, or together with further doses, produces the desired response. In the case of treating a particular disease, such as cancer, the desired response is inhibiting the progression of the disease. This may involve only slowing the progression of the disease temporarily, although more preferably, it involves halting the progression of the disease permanently. This can be monitored by routine methods or can be monitored according to diagnostic methods of the invention discussed herein.

Such amounts will depend, of course, on the particular condition being treated, the severity of the condition, the individual patient parameters including age, physical condition, size and weight, the duration of the treatment, the nature of concurrent therapy (if any), the specific route of administration and like factors within the knowledge and expertise of the health practitioner. These factors are well known to those of ordinary skill in the art and can be addressed with no more than routine experimentation. It is generally preferred that a maximum dose of the individual components or combinations thereof be used, that is, the highest safe dose according to sound medical judgment. It will be understood by those of ordinary skill in the art, however, that a patient may insist upon a lower dose or tolerable dose for medical reasons, psychological reasons or for virtually any other reasons.

The pharmaceutical compositions used in the foregoing methods preferably are sterile and contain an effective amount of for example, a dominant negative

iASPP C6 or nucleic acid encoding a dominant negative iASPP C6, for producing the desired response in a unit of weight or volume suitable for administration to a patient. The response can, for example, be measured by determining the signal transduction inhibited by the dominant negative iASPP

5 C6, composition via a reporter system, by measuring downstream effects such as gene expression, or by measuring the physiological effects of the iASPP C6 composition, such as regression of a tumour, decrease of disease symptoms, modulation of apoptosis, etc.

10 The doses of dominant negative iASPP C6 polypeptide or nucleic acid administered to a subject can be chosen in accordance with different parameters, in particular in accordance with the mode of administration used and the state of the subject. Other factors include the desired period of treatment. In the event that a response in a subject is insufficient at the initial
15 doses applied, higher doses (or effectively higher doses by a different, more localized delivery route) may be employed to the extent that patient tolerance permits.

20 In general, doses of dominant negative iASPP C6 are formulated and administered in doses between 1 ng and about 500mg, and between 10 ng and 100mg, according to any standard procedure in the art. Where nucleic acids encoding dominant negative iASPP C6 are employed, doses of between 1 ng and 0.1mg generally will be formulated and administered according to standard procedures. Other protocols for the administration of iASPP C6 compositions
25 will be known to one of ordinary skill in the art, in which the dose amount, schedule of injections, sites of injections, mode of administration (e.g., intratumoral) and the like vary from the foregoing. Administration of iASPP C6 compositions to mammals other than humans, e.g. for testing purposes or veterinary therapeutic purposes, is carried out under substantially the same
30 conditions as described above. A subject, as used herein, is a mammal,

preferably a human, and including a non-human primate, cow, horse, pig, sheep, goat, dog, cat or rodent.

When administered, the pharmaceutical preparations of the invention are applied in pharmaceutically-acceptable amounts and in pharmaceutically-acceptable compositions. The term "pharmaceutically acceptable" means a non-toxic material that does not interfere with the effectiveness of the biological activity of the active ingredients. Such preparations may routinely contain salts, buffering agents, preservatives, compatible carriers, and optionally other therapeutic agents. When used in medicine, the salts should be pharmaceutically acceptable, but non-pharmaceutically acceptable salts may conveniently be used to prepare pharmaceutically-acceptable salts thereof and are not excluded from the scope of the invention. Such pharmacologically and pharmaceutically-acceptable salts include, but are not limited to, those prepared from the following acids: hydrochloric, hydrobromic, sulfuric, nitric, phosphoric, maleic, acetic, salicylic, citric, formic, malonic, succinic, and the like. Also, pharmaceutically-acceptable salts can be prepared as alkaline metal or alkaline earth salts, such as sodium, potassium or calcium salts.

iASPP C6 compositions may be combined, if desired, with a pharmaceutically-acceptable carrier. The term "pharmaceutically-acceptable carrier" as used herein means one or more compatible solid or liquid fillers, diluents or encapsulating substances which are suitable for administration into a human. The term "carrier" denotes an organic or inorganic ingredient, natural or synthetic, with which the active ingredient is combined to facilitate the application. The components of the pharmaceutical compositions also are capable of being co-mingled with the molecules of the present invention, and with each other, in a manner such that there is no interaction which would substantially impair the desired pharmaceutical efficacy.

The pharmaceutical compositions may contain suitable buffering agents, including: acetic acid in a salt; citric acid in a salt; boric acid in a salt; and phosphoric acid in a salt.

5 The pharmaceutical compositions also may contain, optionally, suitable preservatives, such as: benzalkonium chloride; chlorobutanol; parabens and thimerosal.

10 The pharmaceutical compositions may conveniently be presented in unit dosage form and may be prepared by any of the methods well-known in the art of pharmacy. All methods include the step of bringing the active agent into association with a carrier which constitutes one or more accessory ingredients. In general, the compositions are prepared by uniformly and intimately bringing the active compound into association with a liquid carrier, a finely divided 15 solid carrier, or both, and then, if necessary, shaping the product.

20 Compositions suitable for oral administration may be presented as discrete units, such as capsules, tablets, lozenges, each containing a predetermined amount of the active compound. Other compositions include suspensions in aqueous liquids or non-aqueous liquids such as a syrup, elixir or an emulsion.

25 Compositions suitable for parenteral administration conveniently comprise a sterile aqueous or non-aqueous preparation of iASPP C6 polypeptides or nucleic acids, which is preferably isotonic with the blood of the recipient. This preparation may be formulated according to known methods using suitable dispersing or wetting agents and suspending agents. The sterile injectable preparation also may be a sterile injectable solution or suspension in a non-toxic parenterally-acceptable diluent or solvent, for example, as a solution in 1,3-butane diol. Among the acceptable vehicles and solvents that may be 30 employed are water, Ringer's solution, and isotonic sodium chloride solution. In addition, sterile, fixed oils are conventionally employed as a solvent or

suspending medium. For this purpose any bland fixed oil may be employed including synthetic mono- or di-glycerides. In addition, fatty acids such as oleic acid may be used in the preparation of injectables. Carrier formulation suitable for oral, subcutaneous, intravenous, intramuscular, etc. administrations can be 5 found in Remington's Pharmaceutical Sciences, Mack Publishing Co., Easton, PA.

In a preferred embodiment of the invention said nucleic acid molecule is an inhibitory RNA (RNAi) molecule or antisense nucleic acid molecule.

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In a preferred embodiment of the invention said nucleic acid molecule is selected from the group consisting of an antisense molecule or an inhibitory RNA molecule designed with reference to the nucleic acid sequence shown in Figure 1b. Preferably said antisense or inhibitory RNA molecule is designed to 15 that part of said nucleic acid sequence which encodes an amino acid sequence as defined by amino acid residues +1 to +483 as shown in Figure 1a.

As used herein, the term "antisense molecule" or "antisense" describes an oligonucleotide that is an oligoribonucleotide, oligodeoxyribonucleotide, 20 modified oligoribonucleotide, or modified oligodeoxyribonucleotide which hybridises under physiological conditions to DNA comprising a particular gene or to an mRNA transcript of that gene and, thereby, inhibits the transcription of that gene and/or the translation of that mRNA. The antisense molecules are designed so as to interfere with transcription or translation of a target gene 25 upon hybridisation with the target gene or transcript. Those skilled in the art will recognise that the exact length of the antisense oligonucleotide and its degree of complementarity with its target will depend upon the specific target selected, including the sequence of the target and the particular bases which comprise that sequence. It is preferred that the antisense oligonucleotide be 30 constructed and arranged so as to bind selectively with the target under physiological conditions, i.e., to hybridise substantially more to the target

sequence than to any other sequence in the target cell under physiological conditions. Based upon the iASPP 6C nucleic acid sequences provided herein, or upon allelic or homologous genomic and/or cDNA sequences, one of skill in the art can easily choose and synthesise any of a number of appropriate 5 antisense molecules for use in accordance with the present invention. For example, a "gene walk" comprising a series of oligonucleotides of 15-30 nucleotides spanning the length of iASPP 6C nucleic acid can be prepared, followed by testing for inhibition of the corresponding iASPP 6C expression. Optionally, gaps of 5-10 nucleotides can be left between the oligonucleotides to 10 reduce the number of oligonucleotides synthesised and tested.

In order to be sufficiently selective and potent for inhibition, such antisense oligonucleotides should comprise at least 10 and, more preferably, at least 15 consecutive bases which are complementary to the target, although in certain 15 cases modified oligonucleotides as short as 7 bases in length have been used successfully as antisense oligonucleotides (Wagner et al., *Nature Biotechnol.* 14:840-844, 1996). Most preferably, the antisense oligonucleotides comprise a complementary sequence of 20-30 bases. Although oligonucleotides may be chosen which are antisense to any region of the gene or mRNA transcripts, in 20 preferred embodiments the antisense oligonucleotides correspond to N-terminal or 5' upstream sites such as translation initiation, transcription initiation or promoter sites. In addition, 3'-untranslated regions may be targeted. Targeting to mRNA splicing sites has also been used in the art but may be less preferred if alternative mRNA splicing occurs. In addition, the antisense is targeted, 25 preferably, to sites in which mRNA secondary structure is not expected (see, e.g., Sainio et al., *Cell Mol. Neurobiol.* 14(5):439-457, 1994) and at which proteins are not expected to bind. Finally, although iASPP 6C cDNA sequences are disclosed herein, one of ordinary skill in the art may easily derive the genomic DNA corresponding to the cDNAs. Thus, the present invention 30 also provides for antisense oligonucleotides which are complementary to

IASPP 6C genomic DNA. Similarly, antisense to allelic or homologous cDNAs and genomic DNAs are enabled without undue experimentation.

In one set of embodiments, the antisense oligonucleotides of the invention may 5 be composed of "natural" deoxyribonucleotides, ribonucleotides, or any combination thereof. That is, the 5' end of one native nucleotide and the 3' end of another native nucleotide may be covalently linked, as in natural systems, via a phosphodiester internucleoside linkage. These oligonucleotides may be prepared by art recognised methods which may be carried out manually or by 10 an automated synthesiser. They also may be produced recombinantly by vectors.

In a preferred embodiment of the invention there is provided a transcription cassette comprising an nucleic acid sequence operatively linked to a promoter 15 which promoter transcribes said nucleic acid molecule to produce an antisense nucleic acid molecule, said sequence selected from the group consisting of:

- 20 i) a nucleic acid sequence, or part thereof, as represented in Figure 1b;
- ii) a nucleic acid sequence which hybridises to the sense sequence presented in Figure 1b and which encodes a polypeptide according to the invention.

A recent technique to specifically ablate gene function is through the introduction of double stranded RNA, also referred to as inhibitory RNA 25 (RNAi), into a cell which results in the destruction of mRNA complementary to the sequence included in the RNAi molecule. The RNAi molecule comprises two complementary strands of RNA (a sense strand and an antisense strand) annealed to each other to form a double stranded RNA molecule. The RNAi molecule is typically derived from exonic or coding sequence of the 30 gene which is to be ablated.

Recent studies suggest that RNAi molecules ranging from 100-1000bp derived from coding sequence are effective inhibitors of gene expression. Surprisingly, only a few molecules of RNAi are required to block gene expression which implies the mechanism is catalytic. The site of action appears to be nuclear as 5 little if any RNAi is detectable in the cytoplasm of cells indicating that RNAi exerts its effect during mRNA synthesis or processing.

In a further preferred embodiment of the invention there is provided a transcription cassette comprising a nucleic acid molecule, or part thereof, 10 selected from the group consisting of:

- 15 i) a nucleic acid molecule represented by the nucleic acid sequence in Figure 1b;
- ii) a nucleic acid molecule which hybridises to the sequence in (i) above and which encodes a polypeptide according to the invention; or
- 20 iii) a nucleic acid molecule which is degenerate because of the genetic code to the sequences defined in (i) and (ii) above; wherein said cassette is adapted such that both sense and antisense nucleic acid molecules are transcribed from said cassette.

In a preferred embodiment of the invention said cassette is provided with at least two promoters adapted to transcribe both sense and antisense strands of said nucleic acid molecule.

25 In a further preferred embodiment of the invention said cassette comprises a nucleic acid molecule wherein said molecule comprises a first part linked to a second part wherein said first and second parts are complementary over at least part of their sequence and further wherein transcription of said nucleic acid 30 molecule produces an RNA molecule which forms a double stranded region by complementary base pairing of said first and second parts.

In a preferred embodiment of the invention said first and second parts are linked by at least one nucleotide base.

5 In a preferred embodiment of the invention said first and second parts are linked by 2, 3, 4, 5, 6, 7, 8, 9 or at least 10 nucleotide bases.

In a further preferred embodiment of the invention the length of the RNAi molecule is between 100bp-1000bp. More preferably still the length of RNAi
10 is selected from 100bp; 200bp; 300bp; 400bp; 500bp; 600bp; 700bp; 800bp; 900bp; or 1000bp. More preferably still said RNAi is at least 1000bp.

In an alternative preferred embodiment of the invention the RNAi molecule is between 15bp and 25bp, preferably said molecule is 21bp.

15

In a preferred embodiment of the invention said cassette is part of a vector.

According to a further aspect of the invention there is provided a screening method to identify an agent which modulates the interaction of p53 binding
20 proteins with a p53 polypeptide wherein said method comprises the following steps of:

- i) forming a preparation comprising a polypeptide according to the invention and a p53 polypeptide, or sequence variant thereof, and at least one agent to be tested;
- 25 ii) determining the activity of said agent with respect to the binding of said polypeptide to p53 polypeptide.

According to a further aspect of the invention there is provided a screening method for the identification of an agent which modulates the interaction of
30 Bcl-2 binding polypeptides with a Bcl-2 polypeptide wherein said method comprises the steps of:

5

- i) forming a preparation comprising a polypeptide as represented by the amino acid sequence shown in Figure 2a, or a variant polypeptide which is modified by addition deletion or substitution of at least one amino acid residue and a Bcl-2 polypeptide or variant thereof, and at least one agent to be tested; and
- ii) determining the activity of said agent with respect to the binding of said polypeptide for said Bcl-2 polypeptide.

10 According to a yet further aspect of the invention there is provided a screening method to identify agents which modulate the ubiquitination of a polypeptide according to the invention comprising the steps of:

15

- i) forming a preparation comprising a polypeptide according to the invention, a ubiquitin polypeptide or variant thereof, polypeptide(s) with the specific activity associated with ubiquitin conjugating polypeptides and at least one agent to be tested;
- ii) determining the activity of said agent with respect to the conjugation of ubiquitin to said polypeptide.

20

In a preferred method of the invention said agent is a peptide or polypeptide.

25

In a preferred method of the invention said peptide is at least 6 amino acid residues in length. Preferably the length of said peptide/polypeptide is selected from the group consisting of: at least 7 amino acid residues; 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 amino acid residues in length. Alternatively the length of said peptide/polypeptide is at least 20 amino acid residues; 30; 40; 50; 60; 70; 80; 90; or 100 amino acid residues in length.

30 It will be apparent to one skilled in the art that modification to the amino acid sequence of peptides agents could enhance the binding and/or stability of the

peptide with respect to its target sequence. In addition, modification of the peptide may also increase the *in vivo* stability of the peptide thereby reducing the effective amount of peptide necessary to inhibit p53 binding of iASPP. This would advantageously reduce undesirable side effects which may result in vivo. Modifications include, by example and not by way of limitation, acetylation and amidation. Alternatively or preferably, said modification includes the use of modified amino acids in the production of recombinant or synthetic forms of peptides. It will be apparent to one skilled in the art that modified amino acids include, by way of example and not by way of limitation, 4-hydroxyproline, 5-hydroxylysine, N⁶-acetyllysine, N⁶-methyllysine, N⁶,N⁶-dimethyllysine, N⁶,N⁶,N⁶-trimethyllysine, cyclohexylalanine, D-amino acids, ornithine. Other modifications include amino acids with a C₂, C₃ or C₄ alkyl R group optionally substituted by 1, 2 or 3 substituents selected from halo (eg F, Br, I), hydroxy or C₁-C₄ alkoxy. Modifications also include, by example and not by way of limitation, acetylation and amidation.

In a preferred embodiment of the invention said peptide sequence is acetylated. Preferably said acetylation is to the amino terminus of said peptide.

20 In a further preferred embodiment of the invention said peptide sequence is amidated. Preferably said amidation is to the carboxyl-terminus of said peptide.

25 It will also be apparent to one skilled in the art that peptides could be modified by cyclisation. Cyclisation is known in the art, (see Scott *et al* Chem Biol (2001), 8:801-815; Gellerman *et al* J. Peptide Res (2001), 57: 277-291; Dutta *et al* J. Peptide Res (2000), 8: 398-412; Ngoka and Gross J Amer Soc Mass Spec (1999), 10:360-363.

In a further preferred method of the invention said antagonist is an antibody or antibody binding part. Preferably said antibody is a monoclonal antibody or binding part thereof.

5 In an alternative preferred method of the invention said agent is an aptamer.

Nucleic acids have both linear sequence structure and a three dimensional structure which in part is determined by the linear sequence and also the environment in which these molecules are located. Conventional therapeutic molecules are small molecules, for example, peptides, polypeptides, or antibodies, which bind target molecules to produce an agonistic or antagonistic effect. It has become apparent that nucleic acid molecules also have potential with respect to providing agents with the requisite binding properties which may have therapeutic utility. These nucleic acid molecules are typically referred to as aptamers. Aptamers are small, usually stabilised, nucleic acid molecules which comprise a binding domain for a target molecule. A screening method to identify aptamers is described in US 5,270,163 which is incorporated by reference. Aptamers are typically oligonucleotides which may be single stranded oligodeoxynucleotides, oligoribonucleotides, or modified oligodeoxynucleotide or oligoribonucleotides.

The term "modified" encompasses nucleotides with a covalently modified base and/or sugar. For example, modified nucleotides include nucleotides having sugars which are covalently attached to low molecular weight organic groups other than a hydroxyl group at the 3' position and other than a phosphate group at the 5' position. Thus modified nucleotides may also include 2' substituted sugars such as 2'-O-methyl-; 2-O-alkyl; 2-O-allyl; 2'-S-alkyl; 2'-S-allyl; 2'-fluoro-; 2'-halo or 2-azido-ribose, carbocyclic sugar analogues a-anomeric sugars; epimeric sugars such as arabinose, xyloses or lyxoses, pyranose sugars, furanose sugars, and sedoheptulose.

Modified nucleotides are known in the art and include by example and not by way of limitation; alkylated purines and/or pyrimidines; acylated purines and/or pyrimidines; or other heterocycles. These classes of pyrimidines and purines are known in the art and include, pseudoisocytosine; N4, N4-5 ethanocytosine; 8-hydroxy-N6-methyladenine; 4-acetylcytosine, 5-(carboxyhydroxymethyl) uracil; 5-fluorouracil; 5-bromouracil; 5-carboxymethylaminomethyl-2-thiouracil; 5-carboxymethylaminomethyl uracil; dihydrouracil; inosine; N6-isopentyl-adenine; 1-methyladenine; 1-methylpseudouracil; 1-methylguanine; 2,2-dimethylguanine; 2-methyladenine; 10 2-methylguanine; 3-methylcytosine; 5-methylcytosine; N6-methyladenine; 7-methylguanine; 5-methylaminomethyl uracil; 5-methoxy amino methyl-2-thiouracil; β -D-mannosylqueosine; 5-methoxycarbonylmethyluracil; 5-methoxyuracil; 2 methylthio-N6-isopentenyladenine; uracil-5-oxyacetic acid methyl ester; psueouracil; 2-thiocytosine; 5-methyl-2 thiouracil, 2-thiouracil; 4-thiouracil; 5-methyluracil; N-uracil-5-oxyacetic acid methylester; uracil 5-oxyacetic acid; queosine; 2-thiocytosine; 5-propyluracil; 5-propylcytosine; 5-ethyluracil; 5-ethylcytosine; 5-butyluracil; 5-pentyluracil; 5-pentylcytosine; and 2,6,-diaminopurine; methylpsuedouracil; 1-methylguanine; 1-methylcytosine.

20 The aptamers of the invention are synthesized using conventional phosphodiester linked nucleotides and synthesized using standard solid or solution phase synthesis techniques which are known in the art. Linkages between nucleotides may use alternative linking molecules. For example, linking groups of the formula P(O)S, (thioate); P(S)S, (dithioate); P(O)NR'2; 25 P(O)R'; P(O)OR6; CO; or CONR'2 wherein R is H (or a salt) or alkyl (1-12C) and R6 is alkyl (1-9C) is joined to adjacent nucleotides through -O- or -S-. The binding of aptamers to a target polypeptide is readily tested by assays hereindisclosed.

30 An embodiment of the invention will now be described by example only and with reference to the following figures:

Figure 1a is the amino acid sequence of iASPP C6 protein, amino acid sequence underlined is identical to iASPP; Figure 1b is the nucleic acid sequence of iASPP6C;

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Figure 2a is the amino acid sequence of iASPP; Figure 2b is the nucleic acid sequence of iASPP;

Figure 3 is a sequence alignment of full length iASPP6C and iASPP;

10

Figure 4 illustrates the expression of iASPP expression in various cell lines. Full length iASPP 6C is detected at about 100kDa. The samples probed with antibody LX049.3 are: 2.1kb iASPP/RAI (truncated iASPP) in vitro translated; RKO (colon cancer cell line); HeLa; 293 (kidney); MCF7 (breast); SaOS2 (osteosarcoma); HBL100 (breast); H1299 (lung); U937 (lung); U2OS (osteosarcoma);

15

Figure 5 illustrates an immunoprecipitation/western blot using two different iASPP antibodies: LX049.3 is a mouse monoclonal while pAb 18 is a rabbit antibody (epitopes are given in the peptide alignment, see Figure 3);

20

Figure 6a and 6b illustrates that iASPP 6C is ubiquitinated. This process results in the generation of the 83kDa fragment, which is abolished in the presence of MG132 (ubiquitin-proteosome inhibitor). Ubiquitination also appears to be dependent on cell density. The cells were split according the cell density required and MG132 was added the next day (16-24 hrs later).

25

Figure 7 illustrates that p53 preferentially binds to full length iASPP6C, while Bcl-2 preferentially binds iASPP;

30

Figure 8 illustrates the activity of the full length iASPP6C in cells and that iASPP and p53 are involved in the activation of apoptotic genes but not cell cycle regulatory genes, and that it also interacts with p63 and p73; and

5 Figure 9 illustrates that the full length iASPP6C is preferentially expressed in cells rather than iASPP but at different expression levels.

Materials and Methods

10 Standard molecular and cell biology methods have been used in the exemplification of the invention hereindisclosed and can be readily found in published literature. For example, molecular biological techniques can be found in Sambrook et al Molecular Cloning, A Laboratory Manual, Second Edition (Cold Spring Harbor Laboratory Press 1989); immunological
15 techniques used herein can be found in Using Antibodies, A Laboratory Manual, (Cold Spring Harbor Laboratory Press 1999).

20

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Claims

1. An isolated polypeptide wherein said polypeptide is represented by the amino acid sequence as shown in Figure 1a, or a variant polypeptide which 5 variant is modified by addition, deletion or substitution of at least one amino acid residue characterised in that said polypeptide has the following characteristics:
 - i) a polypeptide which preferentially binds the tumour suppressor polypeptide p53 to inhibit the pro-apoptotic activity of p53 when compared to a polypeptide, or variant thereof, as 10 represented by the amino acid sequence as shown in Figure 2a;
 - ii) a polypeptide which includes at least one amino acid residue which residue is ubiquitinated; and
 - iii) a polypeptide which comprises an amino-terminal polypeptide domain wherein said domain is represented between amino acid 15 1 and 483 of the amino acid sequence shown in Figure 1a.
2. A polypeptide according to Claim 1 wherein said polypeptide preferentially binds p53 when compared to a polypeptide represented by the 20 amino acid sequence shown in Figure 2a.
3. A polypeptide according to Claim 1 or 2 wherein said polypeptide is modified by addition, deletion or substitution of at least one amino acid residue wherein said modification is between amino acid residues 1 and 483 of the 25 amino acid sequence presented in Figure 1a.
4. A polypeptide according to any of Claims 1-3 wherein said polypeptide comprises the amino acid sequence shown in Figure 1a.
- 30 5. A polypeptide according to any of Claims 1-4 wherein said polypeptide consists of the amino acid sequence shown in Figure 1a.

6. An isolated nucleic acid molecule wherein said nucleic acid molecule encodes a polypeptide according to any of Claims 1-5.

5 7. An isolated nucleic acid molecule according to Claim 6 wherein said nucleic acid molecule is represented by the nucleic acid sequence shown in Figure 1b or a nucleic acid molecule which hybridises to the sequence shown in Figure 1b, under stringent hybridisation conditions, and which encodes a polypeptide according to any of Claims 1-5.

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8. A nucleic acid molecule according to Claim 6 or 7 wherein said nucleic acid molecule consists of the nucleic acid sequence shown in Figure 1b.

9. A nucleic acid molecule according to any of Claims 6-8 wherein said 15 molecule is a cDNA.

10. A nucleic acid molecule according to any of Claims 6-8 wherein said molecule is genomic DNA.

20 11. A vector comprising a nucleic acid molecule according to any of Claims 6-10.

12. A method for the production of the polypeptide according to any of Claims 1-5, comprising the steps:

25 i) providing a cell transformed/transfected with a nucleic acid molecule or vector according to any of Claims 6-11;

ii) growing said cell in conditions conducive to the manufacture of said polypeptide; and

iii) purifying said polypeptide from said cell, or its growth environment.

13. An antibody, or binding fragment thereof, which binds the polypeptide according to any of Claims 1-5 characterised in that said antibody binds said polypeptide between amino acid residues 1 to 483 of the amino acid sequence shown in Figure 1a.

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14. An antibody according to Claim 13 wherein said fragment is a Fab fragment.

15. An antibody fragment according to Claim 14 wherein said antibody is selected from the group consisting of: F(ab')₂, Fab, Fv and Fd fragments; and antibodies comprising CDR3 regions.

16. An antibody, or binding fragment thereof, according to any of Claims 13-15 wherein said antibody is a humanised.

15

17. An antibody, or binding fragment thereof, according to any of Claims 13-15 wherein said antibody is a chimeric antibody.

18. A polypeptide according to any of Claims 1-5 for use as a pharmaceutical.

19. A nucleic acid molecule or vector according to any of Claims 6-11 for use as a pharmaceutical.

25 20. Use according to Claim 19 wherein said nucleic acid molecule is an inhibitory RNA molecule.

21. Use according to Claim 19 wherein said nucleic acid molecule is an antisense nucleic acid molecule.

30

22. Use according to Claim 20 or 21 wherein said nucleic acid molecule is selected from the group consisting of an antisense molecule or an inhibitory RNA molecule designed with reference to the nucleic acid sequence shown in Figure 3, wherein said antisense or inhibitory RNA molecule is designed to that part of said nucleic acid sequence which encodes amino acid residue 1 to 483 defined as shown in Figure 1a.

5 23. Use according to Claim 22 wherein said nucleic acid molecule is provided as a transcription cassette comprising an nucleic acid sequence 10 operatively linked to a promoter which promoter transcribes said nucleic acid molecule to produce an antisense nucleic acid molecule, said sequence selected from the group consisting of:

- 15 i) a nucleic acid sequence, or part thereof, as represented in Figure 1b;
- ii) a nucleic acid sequence which hybridises to the sense sequence presented in Figure 1b and which encodes a polypeptide according any of Claims 1-6.

24. Use according to Claim 22 wherein said nucleic acid molecule is 20 provided as a transcription cassette comprising a nucleic acid molecule, or part thereof, selected from the group consisting of:

- 25 i) a nucleic acid molecule represented by the nucleic acid sequence in Figure 1b;
- ii) a nucleic acid molecule which hybridises to the sequence in (i) above and which encodes a polypeptide according to any of Claims 1-5; or
- iii) a nucleic acid molecule which is degenerate because of the genetic code to the sequences defined in (i) and (ii) above; wherein said cassette is adapted such that both sense and antisense nucleic acid molecules are transcribed from said cassette.

25. Use according to Claim 24 wherein said cassette is provided with at least two promoters adapted to transcribe both sense and antisense strands of said nucleic acid molecule.

5

26. Use according to Claim 24 wherein said cassette comprises a nucleic acid molecule wherein said molecule comprises a first part linked to a second part wherein said first and second parts are complementary over at least part of their sequence and further wherein transcription of said nucleic acid molecule 10 produces an RNA molecule which forms a double stranded region by complementary base pairing of said first and second parts.

27. Use according to Claim 26 wherein said first and second parts are linked by at least one nucleotide base.

15

28. Use according to any of Claims 23-27 wherein said cassette is part of a vector.

29. A screening method to identify an agent which modulates the 20 interaction of a p53 binding protein with a p53 polypeptide wherein said method comprises the following steps of:

- i) forming a preparation comprising a polypeptide according to any of Claims 1-5 and a p53 polypeptide, or sequence variant thereof, and at least one agent to be tested;
- 25 ii) determining the activity of said agent with respect to the binding of said polypeptide to said p53 polypeptide.

30. A screening method for the identification of an agent which modulates the interaction of a Bcl-2 binding polypeptide with a Bcl-2 polypeptide wherein 30 said method comprises the steps of:

- i) forming a preparation comprising a polypeptide as represented by the amino acid sequence shown in Figure 2a, or a variant polypeptide which is modified by addition deletion or substitution of at least one amino acid residue and a Bcl-2 polypeptide, or variant thereof, and at least one agent to be tested; and
- ii) determining the activity of said agent with respect to the binding of said polypeptide to said Bcl-2 polypeptide.

10 31. A screening method to identify agents which modulate the
ubiquitination of a polypeptide comprising the steps of:

- i) forming a preparation comprising a polypeptide according to any of Claims 1-5, a ubiquitin polypeptide or variant thereof, polypeptide(s) with the specific activity associated with ubiquitin conjugating polypeptides and at least one agent to be tested;
- ii) determining the activity of said agent with respect to the conjugation of ubiquitin to said polypeptide.

20 32. A method according to any of Claims 29-31 wherein said agent is a peptide or polypeptide.

33. A method according to Claim 32 wherein said peptide/polypeptide is an antibody or antibody binding fragment.

25

34. A method according to Claim 31 wherein said agent is an aptamer.

Abstract

Tumour Suppressor Protein

5

We describe a polypeptide which binds and modulates the activity of a tumour suppressor polypeptide, for example p53; a nucleic acid molecule encoding said protein and screening methods which modulate the binding activity of said polypeptide for its target polypeptide(s).

Figure 1a

MDSEAFQSARDFLDMNFQSLAMKHM**DLKQMELDTAAAKVDELT**KOLESLWSDSPAPPQAGPPSRPPRYS
YSSSSIP
EFGSRGSPRKAATDGA**DTPFGRSES**APLHPSPLSPKGRPSSPRTPLYLQPDAYGSLDRATSPRPRAFDGAGSS
LGRAPSRRPGPGLRQQGPPTPFDLGRAGSPRGSPLAEGPQAFFPERGSPRPPATAYDAPASAFGSSLLGSGGS
AFAPPLRAQDDLTLLRRPCKAWNEDLDVAYEKKPSQTASYERLDVFARPASPSLQLLPWRESSLDGLGGTGKDNL
TSATLPRNYKVSPLASDRSDAGSYRRSLGSAGPSTLPRSWQPVSRIPMPSSPQPRGAPRQRPPIPLSMIFKLQN
AFWEHGSRAMLPGSPLFTRAPPKLIQQPQQPQQPQQLPQQTQPTPAQHPQQTWPPVNEGPPK
PTELEPPEPEIEGLLTPVLEAGDVDEGPVARPLSPTLQALPPEAQSVPPELEVARVLAEPRLKRRGSMEQAPA
VALPPTHKKOYOQIISRLFHRHGGPGGGPEPELSPITEGSEARAGPPAPAPPAPIPPPAPSQSSPPEOPOSMEMR
SVLRKAGSPRKRARRALNPLVLLDAALTGELEVVOQAVKEMNDPSOPNEEGITALHNAICGANYSIVDFLITAGA
NVNSPDHSGWPLHCAACNDTVICMIALVOHGAIAFATLSDGATAFEKCDPYREGYADCATYLADVEOSMGLMNS
GAVYALWDYSAEFGDELSFREGESVTVLRRDGPETDWWWAALHGOEYVPRNYFGLFPRVKPORSKV*

Underlined is amino acid sequence identical to iASPP

Figure 1b

CCACCGCGTCCGGGAAGCCCCCAGGTGCCAGGATCTGCCGGGATCCGCCGCCCCGCTCCGGCCGGCACCATGGACAGCG
AGGCATTCCAGAGCGCGGGACTTCTGGACATGAACCTCCAGTCGCTGGCATGAAACACATGGATCTGAAGCA
GATGGAGCTGGACACGGCGGGCAAGGTGGATGAACGTGACCAAGCAGCTGGAGTCGCTGTGGTCAGACTCTCCC
GCCGCCCTCTGGCCCGCAGGCCGGACCCCTTCTAGGCCGCCGGTACAGCTCCAGCTCGATCCCTGAGCCCTTCG
GCAGCCGAGGGTCCCCCGGAAGGGCGGCCACGGCAGACACCCCGTTCGACGATCAGAGAGTGCCCCAAC
CCTACACCCCTACAGCCCCTGTCCTCCCAAGGGACGCCGCTCGCAGCACCCCGCTCACTCTGAGCCGGAC
GCCCTACGGCAGCCTGGACCGCGCACCTGCCGGCGCCCTTCGATGGCGCAGGCAGCTCCCTGGCCGTG
CGGCCCTCCCCCGGGCCGGCCAGGCCGCTCCGCCAGAGGGTCCCCCACGCCCTTCGACTTCTGGCCGCG
AGGCTCCCCCGCGCAGCCCCCTGGCGAGGGGCCAGGCCCTTCGAGCTGGCCGTCACCGCCGCCCC
CCTGCCACGCCCTACGACGCCGAGCTCGCCCTTCGGAGCTCCCTGCTAGGCTCCGGCGCAGCGCATTGCCCC
CGCCCTCTGCGCGCAGACGACCTGACGCTGCGCCGGGGCCCTCCGAAGAGCTGGAACGAGCTGACCTGGACGT
GGCGTACGAGAAGACCTTCGAGACAGCAGAGCTATGAACGCCGCTTCGACGCTTCGCAAGGCCCTGCCGAGC
CTGCAGCTTGGCTTGGAGGGAGAGCAGCCCTGGATGGACTGGGGGCCACCGCAAGGACAACCTCACTAGCGCCA
CCCTGCCGCGAATTACAAGGTCTCTCCCTGGCCAGCAGCCGGCTTCAGACGCCGGCAGCTACCGCGCTCGCTGG
GCTCCCGGGGGCCGCTGGCAGCTTGCCTCGCAGCTGGCAGCCCGTCAAGCCGATCCCATGCCCGCTCAGGCC
CCAGCCCCGGGGGGCCCCCGCGCAGCGCTCCATCCCCCTCAAGCATGATCTCAAGCTGCAAGACGCCCTCTGGAG
CACGGGGCCAGCCGCGCATGCTCCCTGGTCCCCCTCTTCAGCCGAGCAGCCCGCTCAAGCTGCAAGGCCAAC
CACAACCACAGCCCCAGCCACAATACAACACAGGCCAGCTGGGAGGCCACAGGCCAACCCCCAACCCCC
TACCCAGCCCCCAGCATCCCCAACAGACATGGCCCCCTGTGAACGAAGGACCCCCAACCCCCAACGGAGCTG
GAGCCTGAGCCGGAGATAGAGGGCTGCTGACACCAGTGTGAGGCTGGCGATGTGGATGAAGGGCTGTAGCAA
GGCCTCTCAGCCCCACGGGGCTGAGCCAGCAGCCACTGCCACGGGAGGCACAGTGGTGGCCGAGCTGGAGGGAGTGGC
ACGGGTGTTGGCGGAAATTCCCCGGGGCTCAACACGGAGGGCTCATGGAGCAGCCCTGCTGTGGCCCTGCC
CCTACCCACAAGAACAGTACCAAGCAGATCATGCCCTCTTCATCGTCACTGGGGCCAGGGCCAGGGGGGG
CGGAGGCCAGAGCTGTCCCCCATCACTGAGGGATCTGAGGCCAGGGCAGGGGCCCCCTGCTCCCTGCCCCACAGCTCC
CATCCACCCCGGGCCCGCTCCAGAGCAGCCCACCAAGAGCAGCCGAGAGCATGGAGATGCGCTCTGTGCTGCC
AAGGGGGCTCCCCCGCGCAAGGCCGCCGCGCGCCCTCAACCCCTGGTGTGCTCTGGACGCCGGCTGACCG
GGGAGCTGGAGGTGGTGAGCAGGGCTGAGAGGGAGATGAACGCCAGGCCAACAGGCCAGAGGGCATCACTG
TTCGACAACGCCATCTGCCGCCAACACTACTCATCTGTTCTCATGCCCTGCTGCTGCTGTGCTGCC
CGACAGCCACGGCTGGACACCCCTGCACTGCCACCGCTCAGCGACGGCGCCACCGCCCTTCGAGAAAGTGC
CACGGCGCTGCAATCTCGCCACCGCTCAGCGACGGCGCCACCGCCCTTCGAGAAAGTGC
GCGAGGGTTATGCTGACTGCCACCTACCTGGCAGACGTCGAGCAGAGTATGGGCTGATGAACAGCG
GGCAGTGTACGCTCTGGACTACAGGCCAGTTCGGGAGCAGCTGCTCTCCGAGGGCGAGT
CGCTACCGTGTGCGAGGGACGGGCCGGAGACCGACTGGTGGGGCCGCTGCACGGCCAG
AGGGCTACGTGCCGCGGAACACTACTCTGGCTGTTCCCAGGGTGAAGCCTCAAAGGAGTAAAGTCTAGC
AGGATAGAAGGAGGTTCTGAGGCTGACAGAAACAAGCATTCTGCTTCCCTCCAGACCTCTCCCTCTG
TTTTTGCTGCCCTTATCTGACCCCTCACCCCTGCTGGTGGTGGTCCCTGCAACGGGTTCTGTTCTGCC
GAACTGCCAGGGAAAGAAGGAGGGGGCCAGGCTTAAATTAGTAATCTGCCCTAGCCTGGGAGGTCTGGG
AAGGGCTGGAAATCACTGGGACAGGAAACCAACTTCTCTTGGCAAAATCAGATCCCGTCAAAGTGCCTC
CCATGCCCTACCAACCATCATCACATCCCCCAGCAAGCCAGCCA
CCACACCACGGATATTCTGGGAGTCAGCTGCTGACACCCATCTCCCAAGCAGCTGGGGCTGGGATGGGCCA
GAAACATTGGCTCTACCAGGATCCCTGCCCAACCTCTCCCAATTAAAGTGCCTCACACAGCTCTGGTTT
AATGTTATAAACAAAATAGAGAAACTTCCCTATAAAATAAAAGTAGTTGCACAGAAAAAA
AA

Figure 2a

iASPP

MWMKDVARPLSPTRLQPALPPEAQSVPELEEVARVLAEIPRPLKRRGSMEQAPAVALPPTHKKQYQQII
SRLFHRHGGPGPGGRSQSCPSSLRPGQGPPLLPHQLPFHRPAPSQSSPPEQPQSMEMRSVLRKAGSPR
KARRARLNPLVLLDAALTGELEVQQAVKEMNDPSQPNEEGITALHNAICGANYSIVDFLITAGANVNS
PDSHGWTPLHCAASNDTVICMALVQHAAIFATLSDGATAFEKCDPYREGYADCATYLADVEQSMG
LMNSGAVYALWDYSAEFGDELSFREGESVTLRRDGPETDWWWAALHGQEYVPRNYFGLFPRVKP
QRSKV

Figure 2b

1 GCGGCCGGT CGACCCGGCG TTCAGACGCG GGCAGCTACC GGCGCTCGCT GGGCTCCGCG
61 GGGCCGTCGG GCACTTMGCG TCGCAGCTGG CAGCCCGTCA GCGCATCCC CATGCCCGCC
121 TCCAGCCCCC AGCCCCGGG GGCCCCGCG CAGCGTCCC TCCCCCTCAG CATGATCTTC
181 AAGCTGCAGA ACAGCCTCTG GGAGCACGGG GCGAGCCGG CCATGCTCCC TGGGTCCCCC
241 CTCTTCACCC GAGCACCCCC GCCTAAGCTG CAGCCCCAAC CACAACCACA GCCCCAGCCA
301 CAATCACAAAC CACAGCCCCA GCTGCCCCA CAGCCCCAGA CCCAACCCCCA AACCCCTACC
361 CCAGCCCTCC ACATCCGCAT CCCAACAGA CATGGCCCCC TGTGAACGAA GGACCCCCCA
421 AACCCCCCAC CGAGCTGGAG CCTGAGCCGG AGATAGAGGG GCTGCTGACA CCAGTGTGCTGG
481 AGGCTGGCGA TGTGGATGAA GGACCCCTGA GCAAGGCTC TCAGCCCCAC GAGGCTGCA
541 CCAGCACTGC CACCGGAGGC ACAGTCGGTG CCCGAGCTGG AGGAGGTGGC ACGGGTGTG
601 GCGGAAATTG CCCGGCCCT CAAACGCAGG GGCTCCATGG AGCAGGCCCC TGCTGTGGCC
661 CTGCCCCCTA CCCACAAGAA ACAGTACCAAG CAGATCATCA GCGCCCTCTT CCATCGTCAT
721 GGGGGGCCAG GGCCCCGGGG GCGGAGCCAG AGCTGTCCCC CATCACTGAG GGATCTGAGG
781 CCAGGGCAGG GCCCCCTGCT CCTGCCCCAC CAGCTCCCCT TCCACCGCCC GGCCCCGTCC
841 CAGAGCAGCC CACCAGAGCA GCGCAGAGC ATGGAGATGC GCTCTGTGCT GCGGAAGGGC
901 GGCTCCCCCG GCAAGGCCCC CGCGCGCGC CTCAACCCCTC TGGTGTCCCT CCTGGACGCG
961 GCGCTGACCG GGGAGCTGGA GGTGGTGCAG CAGGCGGTGA AGGAGATGAA CGACCCGAGC
1021 CAGCCCAACG AGGAGGGCAT CACTGCCCTG CACAACGCCA TCTGCGCGC CAACTACTCT
1081 ATCGTGGATT TCCTCATCAC CGCGGGTGCC AATGTCAACT CCCCCGACAG CCACGGCTGG
1141 ACACCCCTGTC ACTGCGCGGC GTCGTGCAAC GACACAGTC TCTGCATGGC GCTGGTGCAG
1201 CACGGCGCTG CAATCTTCG CACCAAGCTC AGCGACGGCG CCACCGCCTT CGAGAAGTGC
1261 GACCCCTTACG GCGAGGGTTA TGCTGACTGC GCCACCTACC TGGCAGACGT CGAGCAGAGT
1321 ATGGGGCTGA TGAACAGCGG GGCAGTGAC GCTCTCTGGG ACTACAGCGC CGAGTTCGGG
1381 GACGAGCTGT CCTTCCCGCA GGGCGAGCTG GTCCACCGTC TGCGGAGGG ACGGGCCGGAG
1441 GAGACCGACT GGTGGTGGGC CGCGCTGCAC GGCCAGGAGG GCTACGTGCC GCGGAACTAC
1501 TTCTGGGCTGT TCCCCAGGGT GAAGCCTCAA AGGAGTAAAG TCTAGCAGGA TAGAAGGAGG
1561 TTCTGAGGC TGACAGAAAC AAGCATTCCT GCCTTCCCTC CAGACCTCTC CCTCTGTTT
1621 TTGCTGCCCT TATCTGCACC CCTCACCCCTG CTGGTGGTGG TCCCTGCCAC CGGTTCTCTG
1681 TTCTCCCTGGA AGTCCAGGGGA AGAAGGAGGG CCCCAGCCTT AAATTAGTA ATCTGCCCTA
1741 GCCTTGGGAG GTCTGGGAG GGCTGGAAAT CACTGGGAC AGGAAACAC TCCCTTTTG
1801 CAAATCAGAT CCCGTCAAA GTGCCCTCCA TGCCTACAC CATCATACAC TCCCCCAGCA
1861 AGCCAGCCAC CTGCCAGCC GGGCTGGGA TGGGCTACCA CACCACTGGA TATTCCCTGGG
1921 AGTCACTGCT GACACCATCT CTCCCAGAG TCTTGGGTC TGGGTGGGAA ACATTGGTCT
1981 CTACCAAGGAT CCCTGCCCA CCTCTCCCCA ATTAAGTGCC TTCACACAGC ACTGGTTAA
2041 TGTTTATAAA CAAAATAGAG AAACCTGGTTT AATGTTATA AACAAAATAG AGAAAACCTTC
2101 GCTTATAAAAT AAAAGTAGTT TGACACAGAAA TGAAAAAAA AAAAAAAA AAAAAA

Formatted Alignments

216

3-2

Fig 3-3

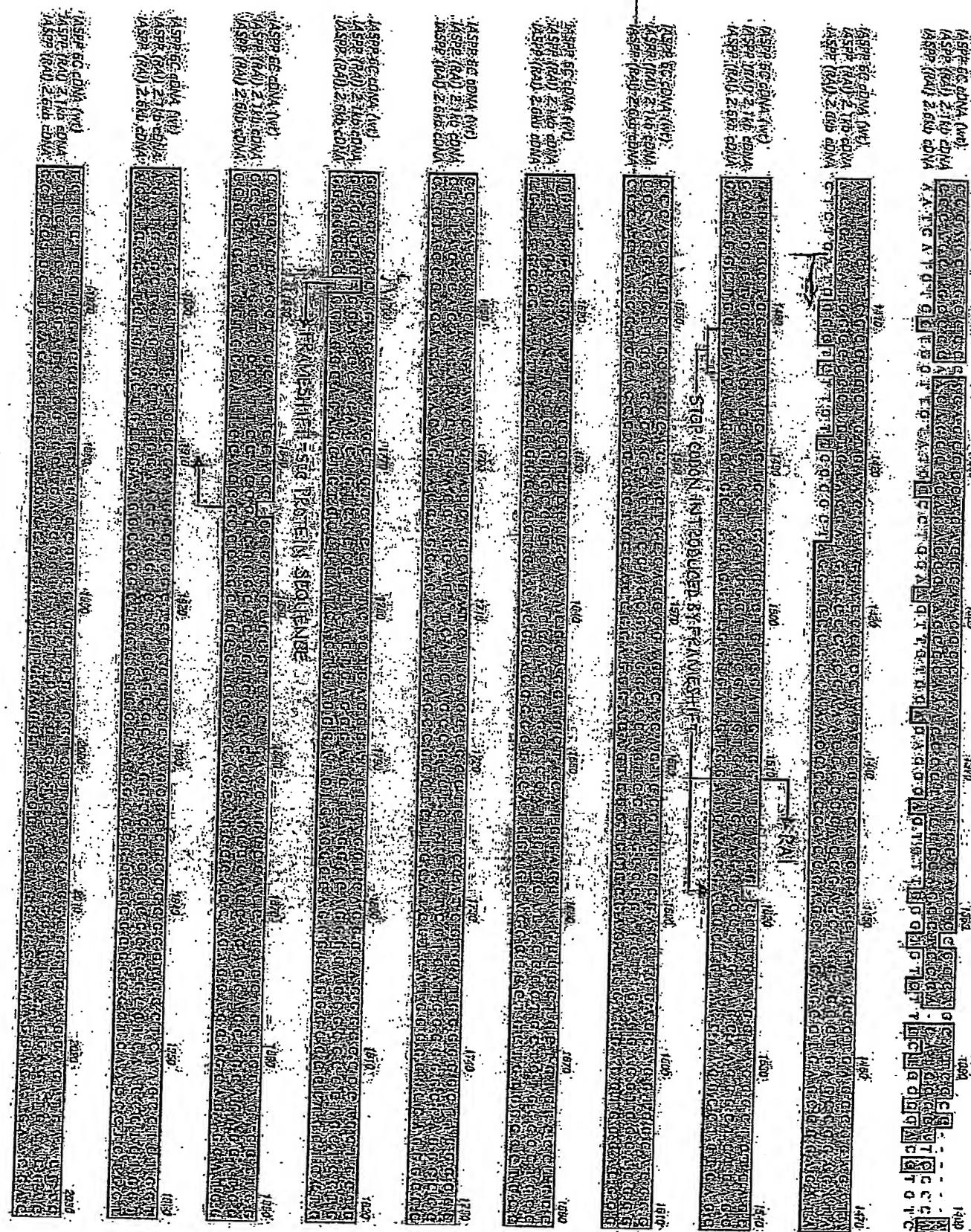


Fig 3-4

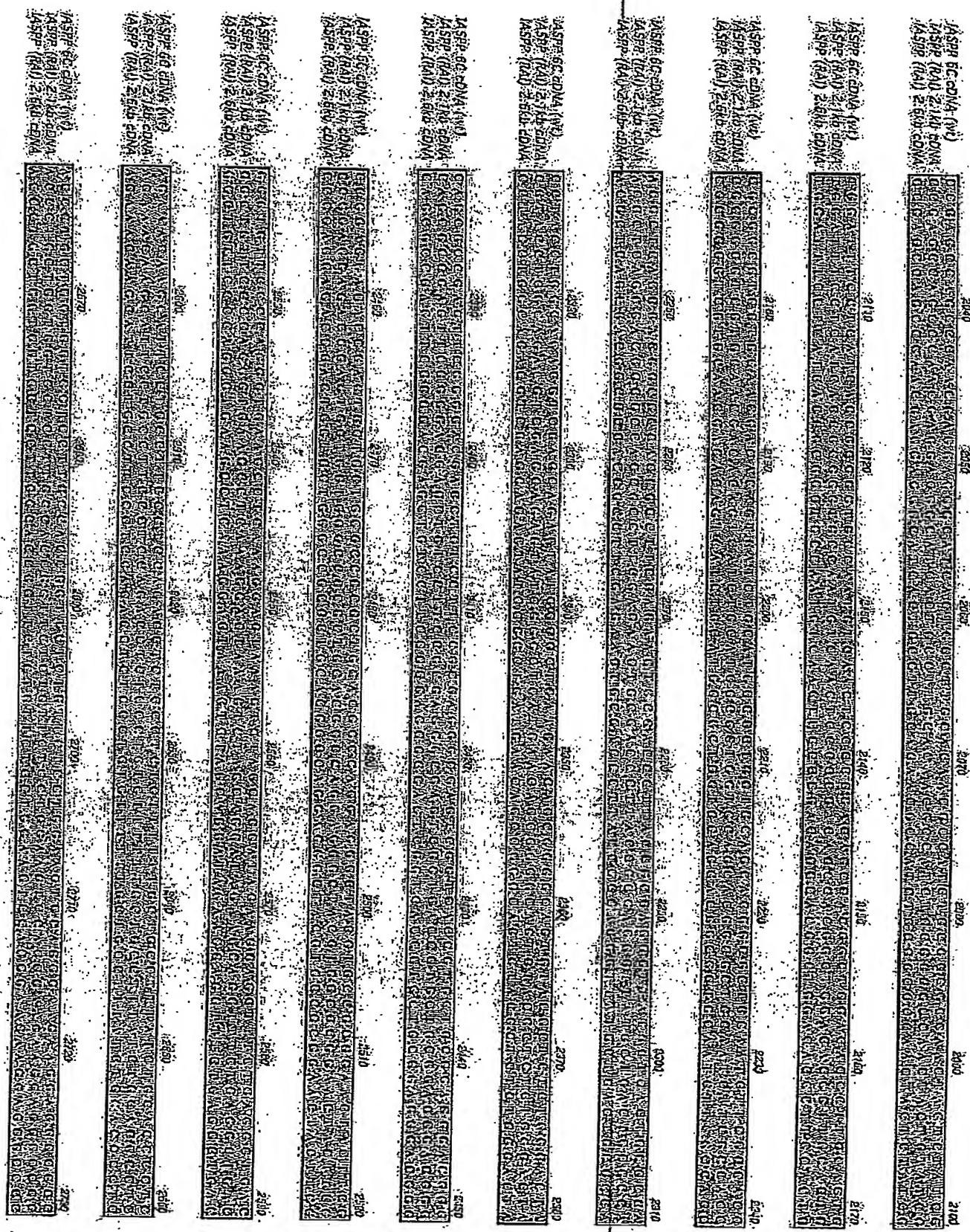


Fig. 3-5.

Formatted Alignment

ASPR/2.5 m connected TIN
ASPR/2.5 m connected TIN
ASPR/2.5 m connected TIN

ASPR/2.5 m connected TIN
ASPR/2.5 m connected TIN
ASPR/2.5 m connected TIN

ASPR/2.5 m connected TIN
ASPR/2.5 m connected TIN
ASPR/2.5 m connected TIN

ASPR/2.5 m connected TIN
ASPR/2.5 m connected TIN
ASPR/2.5 m connected TIN

Fig 3-6

ASPR/2.5 m connected TIN
ASPR/2.5 m connected TIN
ASPR/2.5 m connected TIN

ASPR/2.5 m connected TIN
ASPR/2.5 m connected TIN
ASPR/2.5 m connected TIN

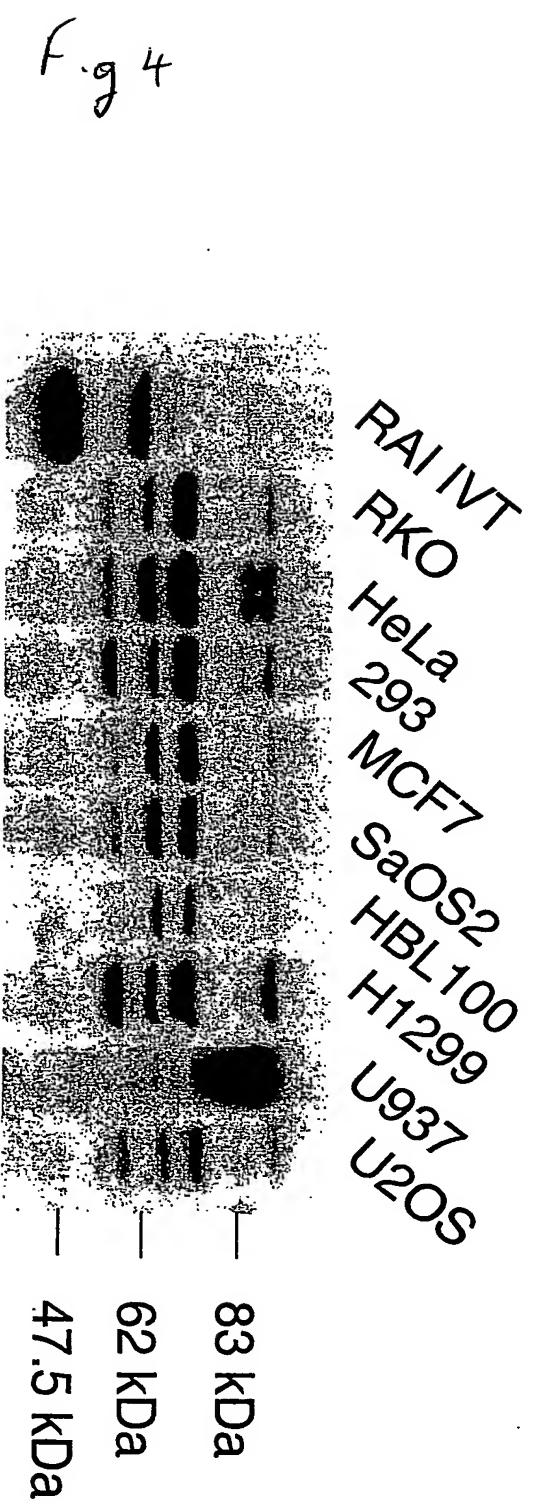
ASPR/2.5 m connected TIN
ASPR/2.5 m connected TIN
ASPR/2.5 m connected TIN

ASPR/2.5 m connected TIN
ASPR/2.5 m connected TIN
ASPR/2.5 m connected TIN

ASPR FRAGMENT USED FOR
LX-093 (NOT FRAMED FOR)

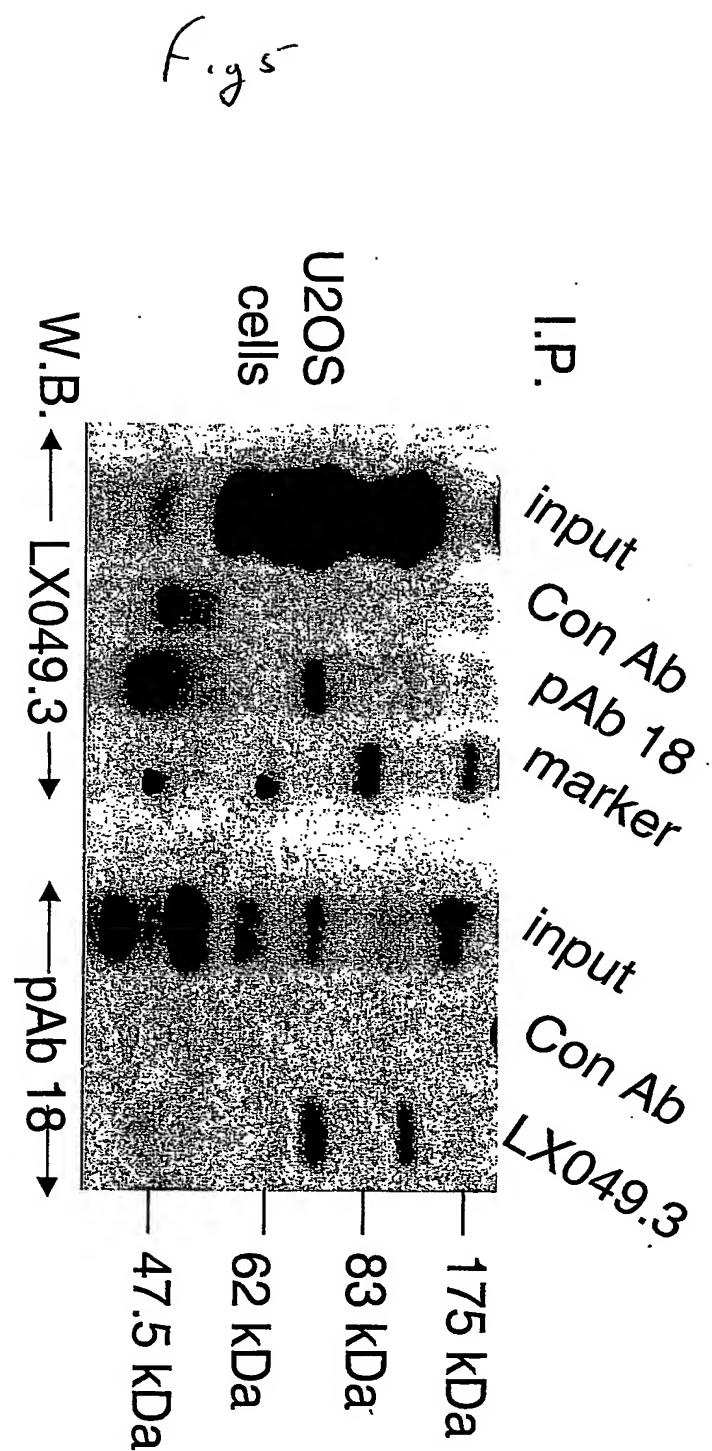
Fig 3-7

Expression of iASPP in various cell lines

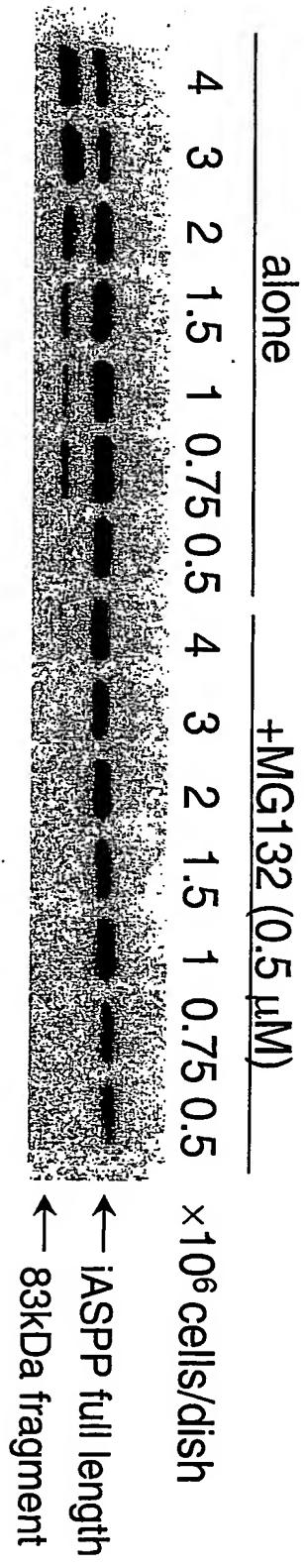


Antibody = LX049.3

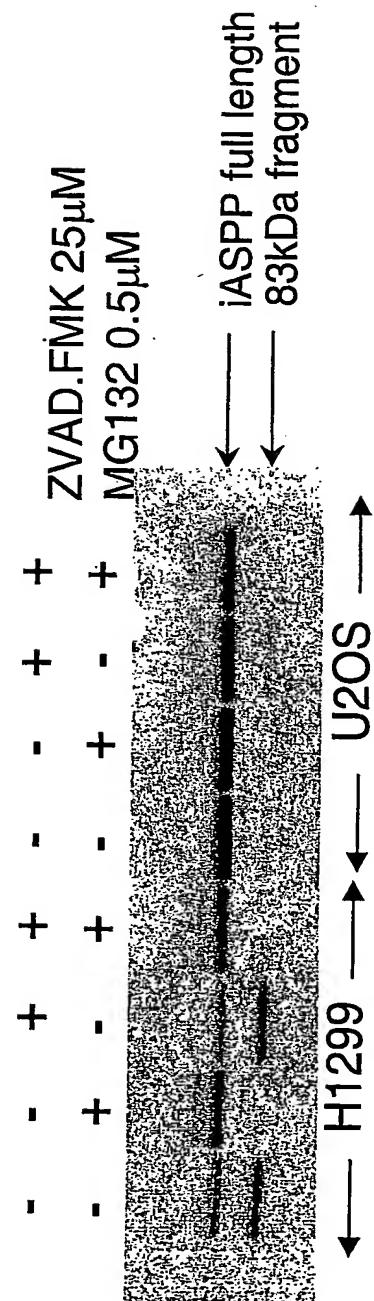
iASPP - I.P./Western blot



Effect of cell density and MG132 upon iASPP expression in U2OS cells



Effect of MG132, Z-VAD.FMK upon iASPP



Interaction of iASPP with p53 and Bcl2 in U2OS cells

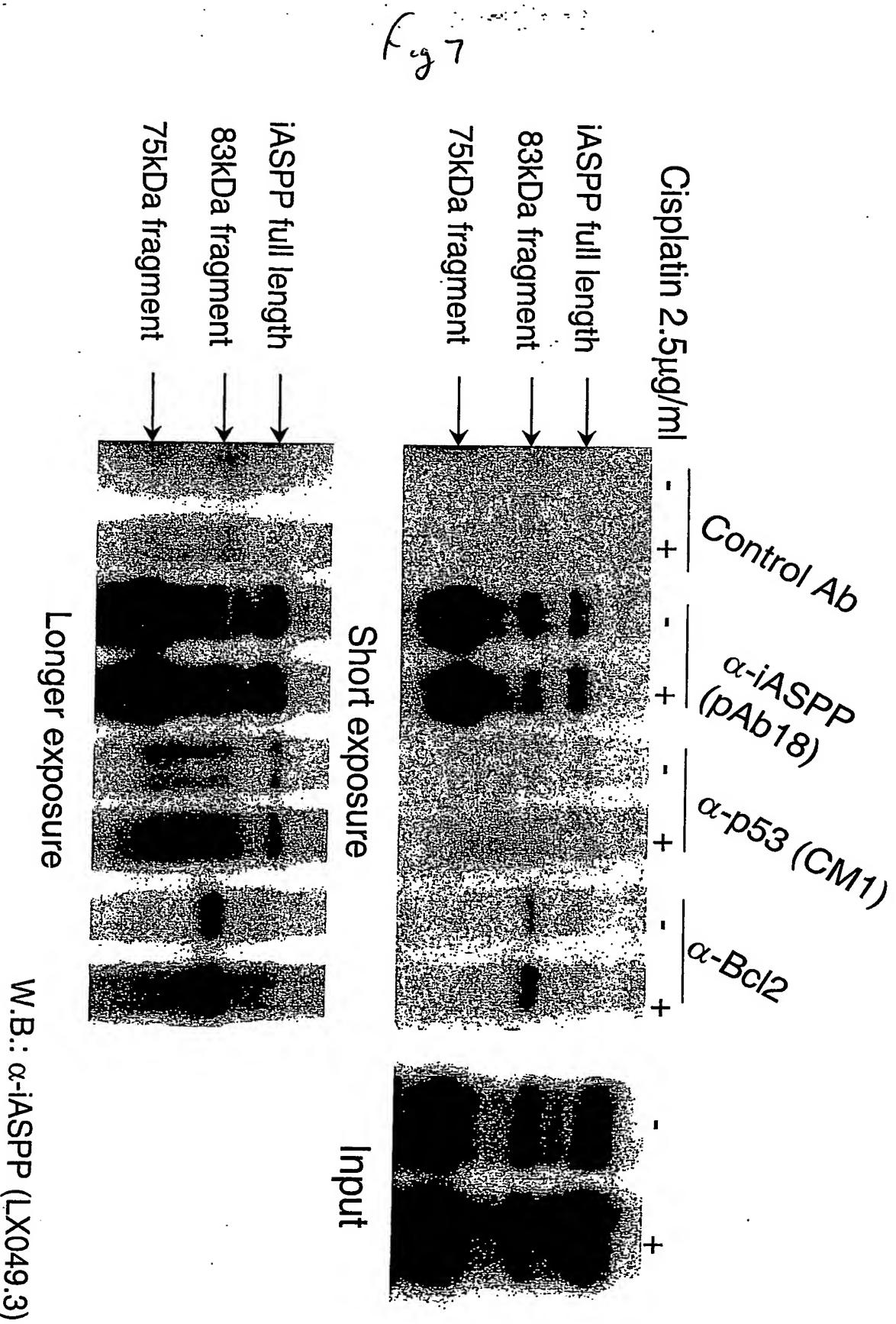
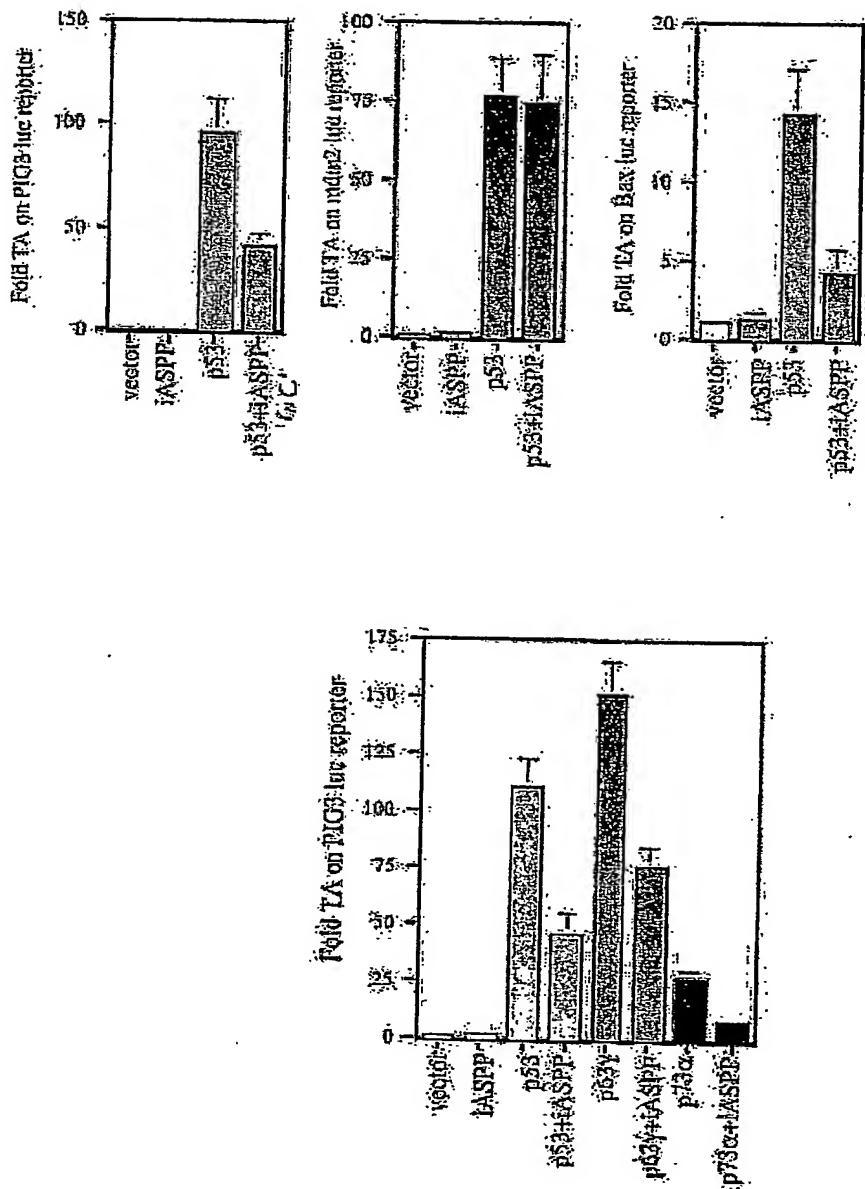
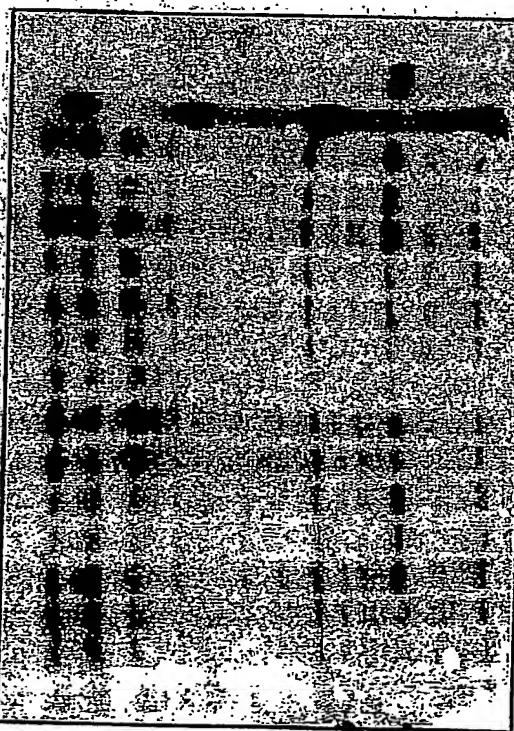
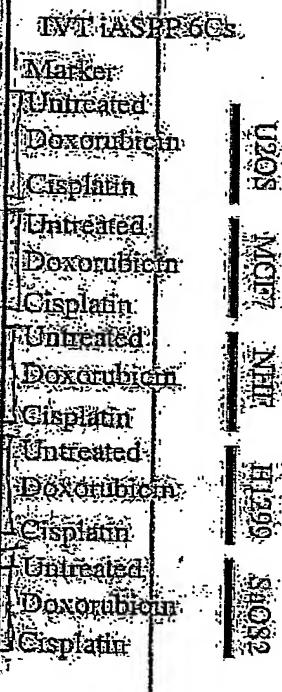


Fig 8



IASPP pattern in five different cell lines



Cell lysates were detected with:
The monoclonal antibody
LX019.3

Cell lysates were detected with:
the monoclonal antibody S442

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